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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Group: 1642 Certificate Under 37 CFR 1.8(a) I hereby certify that this correspondence is being 5816 Confirmation No.: deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to Commissioner for Patents, P. O. Box 1450, Alexandria, VA 22313-1450 Application No.: 09/822,379 Invention: Method of Treatment Using Ligand-Immunogen Conjugates Applicant: Low et al. Ball Rebecca L. Filed: (Printed Name) March 30, 2001 Attorney Docket; 3220-67883 Examiner: Karen A. Canella

DECLARATION UNDER 37 C.F.R. § 1.132 OF DR. BARTON A. KAMEN

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

I declare as follows:

1. I am currently a Professor of Pediatrics and Pharmacology, Director of Pediatric Hernatology-Oncology and Associate Director of the Cancer Institute of New Jersey, University of Medicine and Dentistry of New Jersey. I received a Doctorate of Medicine degree and a Doctorate of Philosophy degree (M.D., Ph.D.) from Case Western Reserve University in 1976. My research has focused on folate and anti-folate metabolism/homeostasis and the function(s) of the folate receptor. I have authored or co-authored more than 150 peer-reviewed journal articles and more than 60 book chapters in the area of my research interests. A copy of my curriculum vitae is attached as Exhibit A.

- 2. I have read and understand the specification of the captioned application and the pending claims in the application. The pending claims of the captioned patent application are directed to methods and compositions for enhancing an endogenous immune response-mediated elimination of a population of cancer cells comprising administering a composition comprising an immunogen conjugated to a folate receptor-binding ligand and a compound capable of stimulating an endogenous immune response wherein the compound does not bind to the conjugate.
- Fig. 1 below. Mice (n = 8 mice/group) were preimmunized with an immunogen and were subsequently injected with M109 cancer cells using an intraperitoneal tumor model. The mice were then injected with a conjugate comprising the immunogen linked to a folate receptor-binding ligand. Control mice were injected with PBS. One group of control mice and one group of the mice treated with the conjugate were then treated with cytokines, compounds capable of stimulating an endogenous immune response. The specific method used is described in detail in Example 7 on page 22 of the patent application. Fig. 1 below is analogous to Fig. 7 in the patent application. The results shown in Fig. 1 below demonstrate that the capacity of a folate-immunogen conjugate to promote long-term survival of tumor-bearing mice is strongly synergistic with cytokines, the cytokines alone having a negligible effect on the survival of the mice in the absence of the folate-immunogen conjugate and the folate-immunogen conjugate alone having only a minor effect.

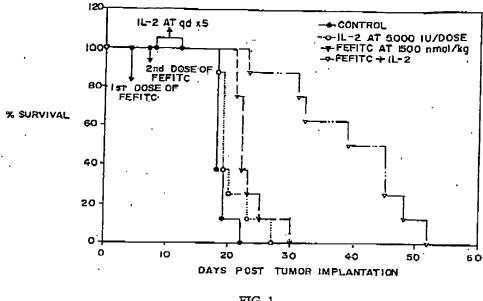
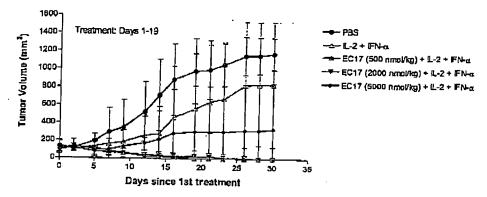


FIG. 1

I understand that the licensee of the captioned application is now in Phase 4. I clinical trials with the method described and claimed in the present application as a cancer therapy. I understand that in vivo assays have been performed by Dr. Yingjuan Lu, a listed inventor on the captioned application and a research scientist for the licensee, for submission to the FDA in Investigational New Drug Study Reports. These assays utilized an M109 subcutaneous tumor model in Balb/c mice, and compared the effect on tumor volume (mm³) of a folate-immunogen conjugate (EC17) in combination with cytokines with the effect on tumor volume of cytokines alone. Control mice were injected with PBS. Exemplary data (n = 8mice/group) generated in such an assay are shown below in Fig. 2. I have studied the data shown in Fig. 2. Tumors were implanted 10 days before treatments were initiated (treatments were initiated on day 1 as shown in Fig. 2).

Subcutaneous M109 tumor



EC17 and L-2 (20,000 LVday) were s.c. dosed at 5 times/weck for 3 weeks; iFN- α (25,000 UVday) was s.c. dosed at 3 times/week for 3-weeks; N = 8

FIG. 2

- 5. The data in Fig. 2 show that in animals treated with the compositions and methods described and claimed in the present application, a complete response (i.e., disappearance of the tumor) was observed in up to 100% of the mice. In contrast, in animals treated with cytokines alone, a maximum response of only 25% was observed. I also understand that in vivo assays have been performed by Dr. Yingjuan Lu which utilized the same tumor model as used for the assay shown in Fig. 2 and the maximum response observed with EC17 alone was 37.5%.
- 6. I understand that in the assay shown in Fig. 2, some of the mice that showed a complete response, in the groups treated with EC17 and cytokines, were subsequently used in other assays. All of the remaining mice that showed a complete response, in the groups treated with EC17 and cytokines, were left untreated for a period of 11 months following tumor implantation and no recurrence of disease was observed. These mice were sacrificed after 11 months to make room for other animals in the animal facility.
- 7. I understand that the effect (i.e., complete tumor disappearance in up to 100% of mice) demonstrated in Fig. 2 is being consistently obtained. These results demonstrate that a complete response (i.e., complete disappearance of tumors) can be obtained in mice with

solid tumors that are treated with folate-immunogen conjugates in combination with cytokines. Cyokines and folate-immunogen conjugates alone each have a considerably reduced effect. A complete response (i.e., complete tumor disappearance) is an unexpected result in the field of cancer therapies utilizing combinations of cancer drugs, particularly where the drugs have never been previously combined.

All statements herein made of my own knowledge are true, and all statements herein made on information and belief are believed to be true; these statements are made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code; and that such willful false statements may jeopardize the validity of the application, any patent issuing thereon, or any patent to which this verified statement is directed.

Dated:

4/13/05

Barton A. Kamen, M.D., Ph.D.

INDS02 RVB 670508+1

CURRICULUM VITAE

Name:

Barton A. Kamen

Birth date/place:

September 30, 1948, New York, NY

Citizenship:

U.S. Citizen

Address:

3 Sleepy Hollow Lane

Princeton Junction, NJ 08550

Phone:

Home: (609) 936-0660

Work: (732) 235-8864 FAX (732) 235-8234

E-mail kamenba@umdnj.edu

Marital Status:

Married 1976, Ruth Saletsky Kamen, Ph.D.

Daughter:

Libby, born 9/24/91

EDUCATION

College:

Western Reserve University, Cleveland Ohio, 1970, B.A.

Graduate School:

Case Western Reserve University, Cleveland, Ohio, 1976,

Ph.D.

Case Western Reserve University, Cleveland, Ohio, 1976.

M.D.

HONORS

Herbert S. Steuer Award for Research in Department of Anatomy,

1976

Alpha Omega Alpha, 1976

Cooley's Anemia Foundation Award (Connecticut), 1978
Damon Runyon/Walter Winchell Cancer Fellowship, 1979-80

Leukemia Society of America Scholar, 1983-87

Burroughs Wellcome Clinical Pharmacology Award, 1987-1992 Founder's Award, Southern Society Pediatric Research, 1989

Student Clinical Teaching Excellence Award, 1991

American Society of Clinical Investigation, 1992

Carl B. and Florence E. King Distinguished Chair, 1993-

American Cancer Society Clinical Research Professorship, 1994-

BOARD CERTIFICATION/MEMBERSHIPS

American Board of Pediatrics
American Board of Pediatrics (Sub-Board of Hematology/Oncology)
American Association for Cancer Research
Society for Pediatric Research
American Federation of Clinical Research
American Society of Clinical Oncology
Southern Society for Pediatric Research
American Society of Clinical Investigation
American Society Pharmacology and Experimental Therapeutics
American Pediatric Society

PROFESSIONAL TRAINING/POSITIONS

1976-77	PL1, Yale University School of Medicine (Pediatric Resident)		
1977-78	PL2, Yale University School of Medicine (Pediatric Resident)		
1978-79	Clinical Fellow Pediatrics, Fellow in Medical Oncology		
19 79-80	Damon Runyon/Walter Winchell Cancer Fellow,		
	Department of Pharmacology and Pediatrics, Yale University		
1980-83	Assistant Professor of Pediatrics and Pharmacology, Midwest		
	Children's Cancer Center, Medical College of Wisconsin,		
	Milwaukee Children's Hospital, Milwaukee, Wisconsin		
1982-83	Member, Graduate School Faculty		
1983-84 Associate Professor of Pediatrics and Pharmacology, I			
·	Children's Cancer Center, Medical College of Wisconsin,		
	Milwaukee Children's Hospital, Milwaukee, Wisconsin		
1984-89	Associate Professor of Pediatrics and Pharmacology, The		
	University of Texas Southwestern Medical School, Dallas, Texas.		
1990-99	Professor of Pediatrics and Pharmacology, The University of Texas		
	Southwestern Medical Center, Dallas, Texas.		
1993-99 Carl B. and Florence E. King Foundation Distinguished			
	Pediatric Oncology Research		
1994-	American Cancer Society Clinical Research Professor		
1999- Professor of Pediatrics and Pharmacology, Director of			
	Hematology-Oncology and Associate Director of the Cancer		
	Institute of New Jersey, University of Medicine and Dentistry of		
	New Jersey Robert Wood Johnson Medical School		
1999-	Member Graduate Program in Cellular and Molecular		
	Pharmacology, Robert Wood Johnson Medical School		
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COMMITTEES\EDITORIAL RESPONSIBILITIES

1981-	Pediatric Oncology Group (POG) New Agent and Pharmacology Committee			
1981-89	Pediatric Oncology Group Lymphoid Disease Committee			
1981-84	Chairman Carcinogen Committee The Medical College of			
	Wisconsin			
1982-84	Member, Graduate Faculty The Medical College of Wisconsin			
1982-83	Vice Chairman, New Agent and Pharmacology Committee Head of New Agent Section, Pediatric Oncology Group			
1983-86	Executive Committee of Hematology/Oncology Discipline Committee Pediatric Oncology Group			
1983-87	Associate Editor, Methotrexate Update			
1903-07 1983-88	Chairman, New Agent and Pharmacology Committee, Pediatric			
1903-00	Oncology Group			
1983-	Advisory Board, Who's Who in America (for Cancer Specialists)			
1984-	Biohazards Committee			
1984-96	Pharmacy & Therapeutics Committee, Children's Medical Center & Parkland Memorial Hospital			
1984-86	Transfusion Committee, Children's Medical Center			
1985-	Head, Pharmacy & Therapeutics, Children's Medical Center			
1986-	Director, Intern Selection Committee, Department of Pediatrics, UTSouthwestern Medical Center			
1986-	Institutional Review Board for local American Cancer Society			
1987-	Editorial Board, Cancer Therapy and Control			
1988-	Member Graduate Faculty, UT Southwestern Graduate School of Biomedical Sciences			
1989-	Secretary Medical Staff, Children's Medical Center			
1990-93	Scientific Advisory Board - Chemotherapy and Hematology, American Cancer Society (National)			
1991-	Advisory Board PDQ (NCI data base for cancer treatment)			
1992-93	Program Planning Committee for the American Society Clinical			
	Oncology for Pediatrics and Pharmacology			
1992-	Editorial Board, Cancer Chemotherapy and Pharmacology			
1993-94	Program Committee, AACR			
1993-	Associate Editor, Cancer Research Therapy and Control			
1993-94	Chairperson/speaker/program committee, AACR symposium,			
1993-	Editorial Board, Pharmacotherapy			
1993-	Review Committee Medical Student Basic Science Education			
1993-	Search Committee, Chair Dept. of Cell Biology and Neuroscience			
1993-	Board of Directors AW Harris Faculty club			

1993-	Elected member-at-large of Medical Service, Research and					
	Development plan at UTSouthwestern					
1994-	Editorial Board, Clinical Cancer Research					
1995-	Education Committee, AACR meeting 1996					
1995-1999	Admissions Committee, MSTP (MD/PhD) program,					
	UTSouthwestem					
1995-1997	Search Committee Director of Simmons Cancer Center					
1996-1999	Editorial Board, Journal of Clinical Oncology					
1996-	Board of Directors: Clayton Dabney Foundation for Kids with					
	Cancer					
1996-	Advisory Group on Research Evaluation and Targeting: American					
	Cancer Society					
1996-	American Society Clinical Oncology Subcommittee Phase I Clinical					
,	Trials					
1996-	Interim Director, Clinical Research Office, Simmons Cancer Center					
1997-	Editorial Board, Pediatric Hematology and Oncology					
1997-	Editorial Board, Cancer Therapeutics					
1998-	Program committee, AACR Phase I clinical trials					
	1998- Advisory Committee Burroughs Wellcome Fund Clinical					
	Scientist Awards					
1998-	Publication Relations and Communications Committee, AACR					
1998-	Scientific Editor, Pharmacotherapy					
1998-	Associate Director Simmons Cancer Center (director of exptl					
therapeutics)					
1998-	Research Program Evaluation Advisory Group, American Cancer					
Society						
1999-	Leukemia and Lymphoma Society of America Translational					
•	Research Grant Review Committee					
2000-	Program Committee AACR and AACR-EORTC joint meeting					
	AACR program committee and session chair: clinical					
	pharmacology, modality based, invited speaker at mini-symposium.					
2001-	Chair, American Cancer Society Nominate Peer Reviewers					
	Advisory Group					
2001-	Editorial Academy, International Journal of Oncology					
2002-	AACR co-chair of discussion session of gender, age and race in					
	clinical trials, pediatric meet the expert and faculty member of					
•	scientist-survivor program (also for 2001 meeting)					
2002-	Editorial Board of CURE					
2002-	Selection Committee AACR fellowships					
2003-	Editorial board, Cancer Therapy					
2003	AACR forum chair/speaker - Metronomics					
2003	AACR faculty/meeting chair scientist-survivor program					
2003	AACR Pediatric Oncology Task Force					
2003	Member Institutional Review Board					
2003	Committee of Review (faculty review)					
2004	AACR co-Chairman, Pediatric Symposium					

2004 Editor-in-Chief Journal of Pediatric Hematology-Oncology
2004 Editorial Board Current Pediatric Reviews
2004 Editorial Board Journal of Chemotherapy
2004 New Jersey Commission on Cancer Research (NJCCR)(one of 11 member board)

Some Examples of Additional Current and Past "Academic" Activities:

PhD Committee(s): University of Buffalo and Roswell Park Memorial Institute Reviewer for: Cancer Research, Clinical Cancer Research, Cancer Chemotherapy Pharmacology, Biochemical Pharmacology, Journal Biological Chemistry, Journal Clinical Investigation, Journal Clinical Oncology, Pediatric Hematology/Oncology, Journal National Cancer Institute, Journal Medical and Pediatric Oncology, Pharmacotherapy, New England Journal of Medicine and others.

Ad Hoc Site Visit Cancer and Pharmacology Programs for NIH/NCI and Experimental Therapeutics study section. Special Reviewer for Scientific Advisory Board of St Judes Children's Research Hospital. Reviewer for NIH contracts/ grants for Phase I/II drugs and the RAID program for new drug development. Reviewer for American Cancer Society, Special Grants. Board of Trustees, Dallas Chapter Leukemia Society of America. Advisory Board, Komen Foundation (breast cancer research). Board member and President elect, Dallas County Chapter American Cancer Society.

Teaching Responsibilities: Texas: In addition to in patient attending 8-10 weeks/year (pediatric oncology), one day/week was spent in clinic with "routine hematology/oncology and ½ day/week seeing adults in consultation for phase I/II therapy. There was also weekly tumor board. I also lecture third year medical students 6 times/yr about pediatric oncology, teach Anti-Cancer Pharmacology to second year medical students and the advanced pharmacology course for graduate students (mechanisms of drug action) and lecture first year medical students about purine and pyrimidine metabolism. New Jersey: Lecture graduate students and medical students about Pediatric Oncology and Pharmacology. Attending physician approximately 8 weeks/year and weekly clininc. First year biochemistry lecture: clinical correlates of purine and pyrimidine metabolism. Participate in Princeton University undergraduate program for pred-medical students.

Past support (not inclusive of industry, local philanthropy and consulting)

American Cancer Society, DHP68 (originally CH228), A Study of Mechanisms of Resistance and Toxicity to Anti-Folates. B Kamen PI, \cong 30% effort. 1981-1996 \cong \$100,000/yr (terminated by age...rules of ACS).

NIH CA34840, Interaction of Organ Irradiation and Cancer Drug Pharmacology. B Kamen, John Holcenberg Co-PI 30%. 4/1/82-3/30/87. Total direct cost \$541,580.

NIH, CA52625-03, Folate receptor as a marker and target in cancer therapy. **B** Kamen, PI, 30% effort. **7/90-6/93**. Total direct costs \$220,000,

NIH, GM43169-01, Molecular cytology of the folate receptor. **Co-PI RGW Anderson, B Kamen**, 5% effort. **8/90-7/95**. Total direct cost \$674,000.

NIH, CA33625, Pediatric Oncology Group Activities. G Buchanan Pl. B Kamen 5%. 1/01/93-12/31/93. Total direct \$189,663.

NIH, CA09640-04, Pediatric Hematology and Oncology research Training Grant. G Buchanan Pl. B Kamen 5%. 03/01/90-02/28/94, Direct (3/93-2/94) \$131,962.

NIH, UO1 CA60431, Pharmacologic Studies of Acute Lymphoblastic Leukemia. B Kamen PI, 10% effort, 9/1/93-8/31/97, Total direct cost \$601,680.

Small grants for therapeutic trials (NIH): A case for aminopterin in the treatment of leukemia. **B Kamen PI**, 5% effort, 6/96-5/31/98. Total direct costs \$101,184.

March of Dimes Birth Defects Foundation: Role of folic acid and homocysteine as risk factors for neural tube defects. R. Finnell Pl. B. Kamen 10%, 6/98-5/2000, total direct ≅ 125,000.

NIH, CA Modulation of drug resistance in CNS tumors. D. Kokkinakis, Pl. B Kamen, 5 % time. 9/98-8/01. Total direct cost \$345,006

Parker Hughes Trust: A case for aminopterin in the treatment of leukemia. B. Kamen PI 1/97-12/97. \$50,000.

Scholar Leukemia Society of America, B. Kamen PI salary support 1983-1987, \$30,000/year.

Burroughs Wellcome Clinical Pharmacology Award, B. Kamen PI salary support 1987-1992, \$50,000/year

Carl B. and Florence E. King Distinguished Chair in Pediatric Oncology Research, B. Kamen PI 1993-1999,

50,000/year.

Texas Dept. of Health. Folate Seroprevalence of Texas Women. B. Kamen, Pl 6/99-01. Total Direct ≅ \$45,000.

HD35396 Folate receptor knockouts, arsenate and birth defects. R. Finnell, Pl, 4/97-3/02. B. Kamen, 5% time. Total direct cost \$789,580.

01-1080-CCR-S-O New Jersey Commission for Cancer Resesarch. Cerebral folate transport:role in Cancer treatment. J. Wollack Pl, B. Kamen 5% time. 6/01-5/03. Total \$97,900, first year \$50,000.

FD-R-001458-NIH, Aminopterin in Adults and Children with Acute Leukemia. B. Karnen, PI 9/30/00-9/29/03. Total direct costs \$116,500.

Current:

American Cancer Society Clinical Research Professorship (CRP-6) B. Kamen, PI Salary Support 1994- \$60,000/ year.

ACS Supplement to above: \$10,000/yr for supplies or partial student stipend.

FD-R-001832, A Pilot Study of Aminopterin for Patients with Acute Lymphoblastic Leukemia. **B. Kamen, PI 9/30/00-9/29/05**. Total direct cost \$440,000.

New Jersey Commission on Cancer Research: Pilot trial of dextromethorphan in cancer related fatigue. S Goodin PI, B. Kamen 5% time. \$35,000 2003-2004

Damon-Runyon Walter Winchell/Eli Lilly Translational Award (mentored research) Pl P. Cole, Mentor B. Kamen. 7/03-6/08. total award \$995,000.

NIH R43 CA109801-01: Novel Oral Trojan-Horse Agents For Osteolytic Cancer Goldstein, Alex, S. PI, B. Kamen, 5% time. 9/04-8/06. Total award \$599,998. First year \$299,999.

PATENTS:

Kamen, B.A. and Zebala, J.A. Therapeutic compositions and methods employing aminopterin. U.S. Patent Pending.

Kamen, B. A. Treatment of antifolate neurotoxicity. U.S. Patent Pending.

PUBLICATIONS:

Rothenberg SP, Frances G, and Kamen BA: Antibodies against folic acid— I. *In vitro* biophysical effect. J Lab Clin Med 74:622-671, 1969.

DaCosta M, Rothenberg SP, and Kamen BA: DNA synthesis in chronic granulocytic leukemic cells containing unsaturated folate binder. Blood 39, 621-627, 1972.

Kamen BA, and Caston JD: Direct radiochemical assay for serum folate: competition between ³H-folic acid and 5-methyltetrahydrofolic acid for a folate binder. J Lab Clin Med 83:164–174, 1974.

Kamen BA and Caston JD: Identification of a folate binder in hog kidney. J Biol Chem 250:2203-2205, 1975.

Kamen BA, Takach PL, Vatev R, and Caston JD: A rapid, radiochemical assay for methotrexate. Anal Biochem 70:54-63, 1976.

Kamen BA and Caston JD: Purification of a folate binding factor in normal human umbilical cord serum. Proc Nat Acad Sci 72:4261-4264, 1975.

Kamen BA and Caston JD: Comments of radiochemical folate assay. Clin Chem 22:1409-1410, 1976.

Kamen BA, Summer CP, and Pearson HA: Exchange transfusion as a treatment for hyperleukocytosis, anemia and metabolic abnormalities in a patient with leukemia. J Pediatr 96:1043-1044, 1980.

Gross S, Kamen BA, Fanaroff A, and Caston JD: Folate compartmentation during developmental maturation. J Pediatr 96:842-844, 1980.

Kamen BA, Cashmore AR, Hsieh P, Dreyer RN, Moroson BA and Bertino JR: Transport of methotrexate in a sensitive and resistant cell line: effect of ³H-methotrexate impurities. J Biol Chem 255:3254-3257, 1980.

Kamen BA, Nylen PA, Camitta BM, and Bertino JR: Methotrexate accumulation in cells as a possible mechanism of chronic toxicity to the drug. Br J Haematol 49:355, 1981.

Krakower GA, Nylen PA, and Kamen BA: Identification and separation of subpicomole amounts of methotrexate polyglutamates in animal and human biopsy material. Anal Biochem 122:412-416, 1982.

Ohnoshi T, Ohnuma T, Takahashi I, Scanlon K, Kamen BA, Holland JF: Establishment of methotrexate-resistant human acute lymphoblastic leukemia cells in culture and effects of folate antagonists. Cancer Res 42:1655-1660, 1982.

Vietti TJ, Steuber CP, Kim TH, Holcenberg J, Kamen B, Murray E, and Capiello V: Mitoxantrone in children with advanced malignant disease: a phase I study. In, New Anticancer Drugs. Raven Press, New York, NY, 1982.

Kamen BA, Holcenberg JS, Siegel SA: Treatment of CNS leukemia with AZQ. Cancer Treatment Report 66:2105-2106, 1982.

Kamen BA, Whyte-Bauer Wand Bertino JR: A mechanism of resistance to Methotrexate. NADPH but not NADH stimulation of methotrexate binding to dihydrofolate reductase. Biochemical Pharmacology 32:1837-1841, 1983.

Harb JM, Werlin SL, Camitta BM, Oechler H, Kamen BA, Blank EL: Hepatic ultrastructure in leukemic children treated with methotrexate and 6-mercaptopurine. Amer J Pediatr Hematol/Oncol 5:323-331, 1983.

Krakower GA and Kamen BA: Comments on pharmacokinetics of erythrocyte and plasma MTX. Cancer Chemother and Pharmacol 10:230, 1983.

Krakower GR, Kamen BA: <u>In situ</u> methotrexate polyglutamate formation in rat tissues. J Pharm Exp Ther 227:633, 1983.

Kamen BA, Eibl B, Cashmore A, Bertino JR: Uptake and efficacy of trimetrexate (TMQ, 2,4, diamino-5 methyl-6-[3,4,5-trimethoxynillnomethyl quinazoline) a non-classical antifolate in MTX resistant leukemia cells in vitro. Biochem Pharmacol 33:1697-1699, 1984.

Kamen BA, Holcenberg JS, Turo K, Whitehead VM: Methotrexate and folate content of erythrocytes in patients receiving oral vs intramuscular therapy with methotrexate. J Pediatr 104:130-133, 1984.

Krakower GR, Kamen BA: The reticulocytic rat: A model for the in <u>situ</u> analysis of methotrexate polyglutamate dynamics. J Pharm Exp Ther 231:43-47, 1984.

Kun LE, Camitta BM, Mulhern RK, Lauer SJ, Kline RW, Casper JT, Kamen BA, Kaplan BM, Barber SW: Treatment of meningeal relapse in childhood acute lymphoblastic leukemia: I. Results of craniospinal irradiation. J Clin Oncol 2:359-364, 1984.

Kamen BA, Holcenberg JS, Moulder JE, Ring BJ, Adams SE, Fish BL: Methotrexate accumulation in rat brain is independent of irradiation and drug schedule. Cancer Res, 44:5092-5094, 1984.

Ohnuma T, Lo RJ, Scanlon K, Kamen BA, Ohnoshi T, Wolman SR, Holland JF: Evolution of methotrexate resistance of human acute lymphoblastic leukemia cells in vitro. Cancer Res 45:1815-1822, 1985.

Civin Cl, Krischer J, Land VJ, Nitschke R, Kamer B, Vats T: Pediatric Oncology Group Phase II trial of AMSA in children with solid tumors. Cancer Treat Report 69:335-336, 1985.

Ungerleider RS, Pratt CB, Vietti TJ, Holcenberg JS, Kamen BA, Glaubiger DL, Cohen LF: Phase I trial of mitoxantrone in children. Cancer Treat Rep 69:403-407, 1985.

Kamen BA, Nylen PA, Whitehead VM, Abelson HT, Dolnick BJ, Peterson DW: Lack of dihydrofolate reductase in human tumor and leukemia cells in vivo. Cancer Drug Delivery 2:133-138, 1985.

Kamen BA, Gunther N, Sowinski N, Rizzo J, Marsik F: Analysis of antibiotic stability in a parenteral nutrition solution. J Infec Dis 4:387-389, 1985.

Zimm S, Ettinger L, Holcenberg J, Kamen BA, Vietti TJ, Belasco J, Shutta N, Balis F, Lavi LL, Collins JM, Poplack DG: Phase I and clinical pharmacologic study of mercaptopurine administered as a prolonged intravenous infusion. Cancer Res 45:1869-1873, 1985.

Kamen BA, Gunther N: Daily unit dosing of antibiotics with a programmable, automated syringe pump. Am J Hosp Pharm 42:2715-2716, 1985.

Winick NJ, Krakower G, Kamen BA: Metabolism of MTX to polyglutamyl derivatives and relationship to folate pools <u>in vivo</u>. Proceedings of Second Workshop on Folyl and Anti-Folyl Polyglutamates. Ed. Goldman ID, Praeger Publ. N.Y. pp 297-307, 1985.

Kamen BA and Caston JD: Properties of a folate binding protein (FBP) isolated from porcine kidney. Blochemical Pharmacol 35:2323-2329, 1986.

Kamen BA, Capdevilla A: Receptor-mediated folate accumulation is regulated by the cellular folate content. Proc Natl Acad Sci USA,83:5983-5987,1986.

Kamen BA, Casper J, Lauer S, Camitta BM, Holcenberg J: Treatment of refractory acute lymphoblastic leukemia with VM-26 and cytosine arabinoside. Cancer Treat Rep 70:935-936, 1986.

Kremer JN, Galivan J, Streckfuss A, Kamen BA: Methotrexate metabolism in blood and liver in rheumatoid arthritis: association with hepatic folate deficiency and formation of polyglutamates. Arth Rheum 29:832-835, 1986.

Moulder JE, Holcenberg JS, Kamen BA, Cheng M, Fish BL: Renal irradiation and the pharmacology and toxicity of methotrexate and cisplatinum. Int J Radiat Oncol Biol Phys 12:1415-1418, 1986.

Pratt CB, Kamen BA, Winick N, Sartain P, Champion JE, Ragab AH, Goren MP: Phase I study of iproplatin in pediatric patients: A Pediatric Oncology Group Study. Cancer Treat Rep 71:87-88, 1987.

Holcenberg JS, Tutsch KD, Earhart RH, Ungerleider RS, Kamen BA, Prett CB, Gribble TJ, Glaubiger DL: Phase I study of ICRF 187 in pediatric cancer patients and comparison of its pharmacokinetics in children and adults. Cancer Treat Rep 70:703-709, 1986.

Kinney TR, Kirscher JP, Starling KA, Kamen BA: Pediatric Oncology Group Phase II trial of MGBG in children with leukemia and lymphoma. Cancer Treat Rep 70:1041-1042, 1986.

Goren MP, Forastiere AA, Wright RK, Horowitz MC, Dodge RK, Kamen BA, Pratt CB: Carboplatin (CBDCA), Iproplatin (CHIP) and high dose cisplatin in hypertonic salt evaluated for tubular nephrotoxicity. Cancer Chemother Pharmacol 19:57-60, 1987.

Winick N, Kamen B, Lester C, Balis F, Poplack D, Holcenberg J: Folate and methotrexate polyglutamate tissue levels in monkeys following low dose methotrexate. Cancer Drug Delivery 4:25-31, 1987.

Holcenberg JS, Moulder JE, Kamen BA, Krailo MD, Fish BL, Ring BJ, Adams S: Chronic effects of fractionated renal irradiation on the pharmacokinetics of intravenous methotrexate. Int J Radiat Oncol Biol Phys 13:759-764, 1987.

Winick NJ, Kamen BA, Streckfuss A, Craig J, McGuirt F, Capizzi RL, Sklar F, Coln D: Methotrexate (MTX) concentration in tumors following low dose MTX. Cancer Chemotherapy & Pharmacology 20:78-80, 1987.

Whitehead VM, Kamen BA, Beaulieu D: Levels of dihydrofolate reductase in livers. Cancer Drug Delivery 4:185-189, 1987.

Kamen BA, Winick NJ: High dose methotrexate: insecure rationale? Biochem Pharmacol 37:2713-2715, 1988.

Kamen BA, Wang MT, Streckfuss AJ, Peryea X, Anderson RGW: Delivery of folates to the cytoplasm of MA104 cells is mediated by a surface membrane receptor that recycles. J Biol Chem 263:13602-13609, 1988.

Lacey SW, Sanders JM, Rothberg KG, Anderson RGW, Kamen BA: Complementary DNA for the folate binding protein correctly predicts anchoring to the membrane by glycosyl-phosphatidylinositol. J Clin Invest 84:715-720, 1989.

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Group:	1642	}	Certificate Under 37 CFR 1.8(2)
Confirmation No.:	5816	}	I hereby certify that this correspondence is being deposited with the United States Postal Service with sufficient postage as first class mail in an envelope
Application No.:	09/822,379	}	addressed to Commissioner for Patents, P. O. Box 1450, Alexandria, VA 22313-1450
Invention;	Method of Treatment Using Ligand-Immunogen Conjugates	} }	m_ 4/22/05
	·-	. }	Rebecco BOL
Applicant:	Low et al.	}	(Signamre)
Filed:	March 30, 2001	}	Rebecca L. Ball (Printed Name)
Attorney Docket:	3220-67883	}	
Examiner:	Karen A. Canella	}	

DECLARATION UNDER 37 C.F.R. § 1.131

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

I, Philip S. Low, a citizen of the United States of America and resident of West Lafayette, Indiana, do declare and say that:

I am one of the named inventors on the captioned application for patent filed on March 30, 2001. The captioned application claims priority to U.S. Patent Application No. 60/193,944, filed on March 31, 2000 and to U.S. Patent Application No. 60/255,846, filed on December 15, 2000. I understand that the Examiner has rejected claims 43, 45, 46, and 50-52 of the application under 35 U.S.C. § 103(a) over Cowan (WO 01/32207) in combination with other references. Cowan was published on May 10, 2001 and has an International Filing Date of January 19, 2000.

The invention described and claimed in the captioned application was conceived and reduced to practice in this country prior to January 19, 2000.

The invention described and claimed in the captioned application comprises methods and compositions for enhancing an endogenous immune response-mediated elimination of a population of cancer cells comprising administering a composition comprising an immunogen conjugated to a folate receptor-binding ligand and a compound capable of stimulating an endogenous immune response wherein the compound does not bind to the conjugate.

Exhibit A is a copy of a figure that contains the same data as is shown in Fig. 1 of the patent application except that for Fig. 1 in the patent application the assay was extended for a longer period of time (i.e., to about 65 days post tumor implantation). The assay from which the data shown in Exhibit A was obtained was performed in my laboratory by Yingjuan Lu, the other named inventor on the captioned application. The date that the assay depicted in Exhibit A was completed is shown on Exhibit A, but that date has been reducted. The reducted date is earlier than January 19, 2000.

The data in the figure shown in Exhibit A was obtained from an assay in which mice were treated with folate-immunogen conjugates in combination with cytokines.

Accordingly, the claimed invention was conceived and reduced to practice in the United States prior to January 19, 2000.

I declare further that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code, and that such willful false statements may jeopardize the validity of the application, or any patent assuing thereon.

Dated:

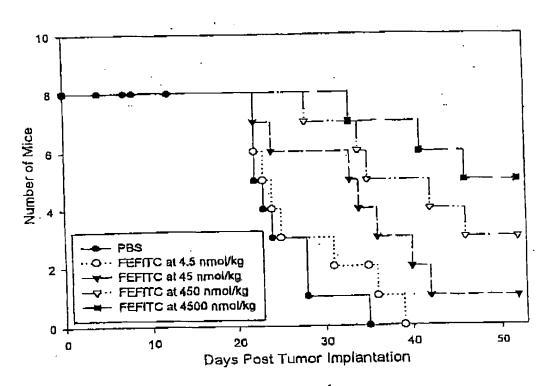
April7, 2005

Ву:

Philip S. Low, Ph.D.

INDS02 RVB 72050640

Fig. 2



*All mice were implanted with i.p. M109 tumors and treated with same doses of IL-2 at a schedule of qd x 5

EXHIBIT A

to 1.131 Declaration